Countermeasures for Treatment

Medication	Administrated for Isotopes	Route of Administration & Dosage	Duration
Aluminum hydroxide ¹	Strontium-90	PO: 60-100 mL	Once
Aluminum phosphate gel ¹	Strontium-90	PO: 100 mL immediately after exposure	Once
Ammonium chloride ^{1 3}	Strontium-90, Radium-226	PO: 1-2 g q.i.d	6 days
Calcium ¹	Strontium-90, Radium-226	PO: Generous doses	
Ca-DTPA ² , Zn-DTPA ²	Plutonium-239, Americium-241, Curium-244, Californium-252 ⁷ , Thorium-232 ⁷ , Yttrium-90 ⁷	1 g in 250 mL NS or 5% glucose, given in 1-2 h, or bolus over 3-4 min Inhalation: 1g in 1:1 dilution with water or NS over 15-20 min IM:	Up to 5 days
Calcium gluconate ¹	Strontium-90,	1 g; not recommended because of pain	6 days
culcium gluconace	Radium-226	5 ampules (500 mg calcium each) in 500 mL D5W over 4 h	o days
Dimercaprol ^{1 4}	Mercury, Lead, Arsenic, Gold, Polonium-210	IM: 300 mg/vial for deep IM use, 2.5 mg/kg (or less) q4h x 2 days, then bid for 1 day, then qd for days 5-10	10 days
D-Penicillamine ¹⁵	Copper, Iron, Mercury, Lead, Gold, and possibly other heavy metals	PO: 250 mg, qd between meals & at bedtime. May increase to 4 or 5 g qd in divided doses.	

Potassium iodide ²	Iodine-131	PO: Adults >40 years of age: with thyroid exposure > 500 cGy: 130 mg/d Adults 18-40 years of age: with thyroid exposure > 50 cGy: 130 mg/d Pregnant or lactating women: 130 mg/d Children and adolescents 3-18: with thyroid exposure > 5 cGy: 65 mg/d Infants 1 month to 3 years: 32.5 mg/d Neonates from birth to 1 month: 16 mg/d (See details)	* In some incidents only a single dose of KI is required. * Incident Managers may recommend additional daily doses if radioactive iodine ingestion (or inhalation) is a continuing threat. * In some incidents, a course of 7-14 days may be recommended.
Potassium phosphate, dibasic ¹	Phosphorus-32	PO: 250 mg phosphorus per tablet. Adult: 1-2 tabs p.o. qid, with full glass of water each time, with meals and at bedtime. Children over 4y: 1 tab qid.	
Propylthiouracil ¹⁶	Iodine-131	PO: 50 mg tabs, 2 tid x 8 days	8 days

Prussian blue ²	Cesium-137, Thallium-201	1 g tid with 100-200 mL water, up to 10 g/d	≥3 weeks, titrated by urine and fecal bioassay and wholebody counting
Sodium alginate ¹	Strontium-90, Radium-226	PO: 10 g powder in a 30 cc vial, add water and drink	
Sodium bicarbonate ¹	Uranium-235	IV: * 2 ampules sodium bicarbonate (44.3 meq each, 7.5%) in 1000 mL NS, 125 mL/h, or * 1 ampule of sodium bicarbonate (44.3 meq, 7.5%) in 500 mL NS, 500 mL/h PO: 2 tablets every 4 h until urine pH=7-8, or 4 g (8 tablets) 3 tid	* Usually IV for the first 24 h, PO for additional 2 days; * Continuation of treatment for >3 days is rare and can be done according to titration of uranium amounts in the body
Sodium phosphate ¹	Phosphorus-32	See Potassium phosphate PO:	3 weeks
Water ¹	Tritium (H-3)	> 3-4 L per day	J WEEKS

NOTES

- 1. Drug listed is not approved by the FDA for this indication / Off label use
- 2. FDA approved for this indication
- 3. FDA Product Safety Information:
 - Oral dosing of ammonium chloride is suggested for the use of radium and strontium internal contamination; however, this indication is not approved, there is currently no approved oral dosage form, and the commercial availability of the oral dosage form is not known.

• Ammonium chloride may be contraindicated in patients with impaired hepatic or renal function.

4. FDA Product Safety Information:

- Adverse events have been reported following use of dimercaprol.
- Use of dimercaprol has been considered in the most severe cases of polonium contamination due to potential toxicity.

5. FDA Product Safety Information:

- Adverse events has been reported following use of penicillamine, including neutropenia, thrombocytopenia, and nephrotoxocity.
- Use of penicillamine has been considered in the most severe cases of cobalt contamination due to potential toxicity.

6. FDA Product Safety Information:

- When KI is unavailable or contraindicated, propylthiouracil and methimazole may be used for blocking thyroid uptake of radioiodine.
- The potential for granulocytopenia and agranulocytosis is significant.
- 7. Ca-DTPA/Zn-DTPA has not been approved by FDA for treating contamination with californium, thorium, and yttrium

